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In the Claims:

1.(currently amended) <u>A recombinant</u> Recombinant cellular system, comprising an animal host cell, comprising the following recombinant proteins:

- a recombinant specific G protein-coupled receptor, and
- <u>a</u> the recombinant <u>CNGA2</u> Ca2+ permeable channel CNGA2.
- 2. (currently amended) The recombinant Recombinant cellular system according to claim 1, <u>further furthermore</u> comprising a recombinant protein <u>selected</u> from the group of connexins, e.g. Cx43 or Cx26.
- 3.(currently amended) The recombinant Recombinant cellular system according to claim 1 or 2, wherein the recombinant specific G protein-coupled receptor is selected from the group of type A guanylyl-cyclases and type G the particular guanylyl-cyclases, e.g. type A to G.
- 4.(currently amended) The recombinant Recombinant cellular system according to claim 1 <u>further</u> or 2, furthermore comprising a cyclase that is harmonised with the specific G protein-coupled receptor, e.g. an adenylyl- or guanylyl-cyclase.
- 5.(currently amended) The recombinant Recombinant cellular system according to claim 1, 2 or 4, wherein the recombinant specific G protein-coupled receptor is selected from: the group of pheromone receptors, e.g. of the V1R-type with all families VR-a to VR-l, including the V3R-type (VR-d), for example V1R-b2, the hormone receptors, e.g. the beta-adrenergic receptors and the olfactory receptors, e.g. OR1A1, OR1A2, Olfr43, Olfr49, MOR261-10, MOR267-1, LOC331758, Olfr41 or Olfr6.

6.(currently amended) The recombinant Recombinant cellular system according to claim 1 further, 2, 4 or 5, furthermore comprising a recombinant G-protein that is harmonised with the specific G protein-coupled receptor, e.g. G-alpha-olf.

7.(currently amended) The recombinant Recombinant cellular system according to claim 1 any of the aforementioned claims, wherein the animal host cell is selected from murine cell lines and or human cell lines, e.g. human cancer cell lines, such as, for example HeLa or HEK293.

8.(currently amended) The recombinant Recombinant cellular system according to claim 1 any of the aforementioned claims, wherein the cellular system comprises a potential recombinant specific G protein-coupled receptor.

9.(currently amended) The recombinant Recombinant cellular system according to claim 7, selected from the group of cellular systems comprising: HeLa-Cx43/CNGA2/Olfr49; HeLa-Cx43/CNGA2/G-alpha-olf; HeLa-Cx43/CNGA2/G-alpha-olf/Olfr41; HeLa-Cx43/CNGA2/G-alpha-olf/Olfr41; HeLa-Cx43/CNGA2/G-alpha-olf/Olfr6 and or HeLa-Cx43/CNGA2/G-alpha-olf/OR1A1.

10.(currently amended) The recombinant Recombinant cellular system according to claim 1 any of the aforementioned claims, wherein the recombinant proteins are present stably and/or transiently transfected.

11.(currently amended) The recombinant —Recombinant cellular system HeLa-Cx43/CNGA2/G-alpha-olf, as deposited on April 20, 2004 at the DSMZ - Deutsche Sammlung von Mikroorganismen and Zellkulturen GmbH in Mascheroder Weg 1b, D-38124 Braunschweig with the deposit number DSM ACC2649.

12.(currently amended) <u>A method</u> for producing a recombinant cellular system, comprising the steps of:

- providing of an animal host cell,
- introducing a recombinant specific G protein-coupled receptor or a potential recombinant specific G protein-coupled receptor, and
- introducing the recombinant <u>CNGA2</u> Ca2+ permeable channel CNGA2.
- 13.(currently amended) <u>The method</u> <u>Method</u> according to claim 12, <u>further</u> <u>furthermore</u> comprising the step of:
- introducing of a recombinant protein from the group of the connexins, e.g. Cx43 or Cx26.
- 14.(currently amended) The method Method according to claim 12 or 13, further furthermore comprising the step of:
- introducing of a cyclase that is harmonised with the specific G protein-coupled receptor, e.g. an adenylyl- or guanylyl-cyclase.
- 15.(currently amended) The method According to claim 12 any of claims 12 to 14, further furthermore comprising the step of:
- introducing of a recombinant G-protein that is harmonised with the specific G protein-coupled receptor, e.g. G-alpha-olf.
- 16.(currently amended) The method Method according to claim 12 any of claims 12 to 15, wherein the introducing method step is selected from:

 (Ca2+-phosphate-)transfection, lipofection or electroporation,

 optionally followed by the step of well as subsequent optional integration into the genome with the aid of a recombinase or and/or antibiotic-selection cloning, or the step of and transduction.
- 17.(currently amended) The method Method for identifying receptor activating substances, comprising the method steps of

- providing a recombinant cellular system according to <u>claim 1</u> any of claims 1 to 7 or 9 to 11,
- contacting of the cellular system with a potential G protein-coupled receptor activating substance, and
- measuring of the activation or inhibition of the Ca2+ influx into the <u>cellular</u> system eell.

18.(currently amended) The method Method according to claim 17, wherein the potential G protein-coupled receptor inducing substance is selected from odorants, such as, for example, (-)citronellal or beta-citronellol, pheromones, and hormones, such as, for example, adrenalin or natriuretic peptide type-C.

19.(currently amended) The method Method according to claim 17 or 18, wherein the measuring of the Ca2+ influx into the cell includes: a loading of the cell with Fura-2-AM or Fluo-4-AM, and measuring of the emission-wavelength at 515 nm.

20.(currently amended) The method Method according to claim 17 any of claims 17 to 19, wherein the cellular system is pre-treated with an enhancer, such as, for example forskolin or thapsigargin.

- 21.(currently amended) <u>A method</u> Method for producing a pharmaceutical composition, comprising the steps of:
- performing a method according to claim 17 any of claims 17 to 20, and
- formulating of the obtained G protein-coupled receptor inducing substance with known auxiliary agents and additives.

22.(currently amended) <u>A method Method</u> for identifying of G protein-coupled receptors, comprising the steps of:

- providing a recombinant cellular system according to claim 8,

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- contacting of the cellular system with a receptor-activating substance or presumably receptor-activating substance, and

- measuring of the activation or inhibition of the Ca2+ influx into the cell.

23.(currently amended) The method Method according to claim 17 any of claims 17 to 22, wherein the method is performed in a high-throughput-environment, e.g. in microtiter-plates in a fluorescence plate reader or high-resolution microscopy supported on the level of individual cells.

24.(cancelled)

25.(cancelled)